10/629,975 Search 6/27/07

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(FILE 'HOME' ENTERED AT 11:55:06 ON 27 JUN 2007)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 11:55:24 ON 27 JUN 2007

	DON 2007	
L1	89	S (IBD TREATMENT)
L2	7	S L1 AND PD<2001
L3	5	DUPLICATE REMOVE L2 (2 DUPLICATES REMOVED)
L4	0	S L3 AND LACTOFER?
L5	0	S L3 AND LEUKOC?
L6	4	S (MONITOR? IBD)
L7	. 2	DUPLICATE REMOVE L6 (2 DUPLICATES REMOVED)

d his

(FILE 'HOME' ENTERED AT 11:55:06 ON 27 JUN 2007)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 11:55:24 ON 27 JUN 2007

L1	89	S	(IBD TREATMENT)			
L2	. 7	S	L1 AND PD<2001			
L3	5	DU	JPLICATE REMOVE L2	(2	DUPLICATES	REMOVED)
L4	0	S	L3 AND LACTOFER?			
L5	0	S	L3 AND LEUKOC?			
L6	4	S	(MONITOR? IBD)			
L7	2	DU	JPLICATE REMOVE L6	(2	DUPLICATES	REMOVED)

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1995:522917 CAPLUS
AN
DN
    122:260562
    Entered STN: 04 May 1995
ED
TI
    Diagnostic test and kit for disease or disorders in the digestive system
    Fagerhol, Magne K.; Dale, Inge; Roseth, Arne G.
IN
PA
    Norway
    Can. Pat. Appl.
SO
    CODEN: CPXXEB
DT
    Patent
LA
    English
IC
    ICM G01N033-68
     ICS G01N033-574
     9-10 (Biochemical Methods)
    Section cross-reference(s): 14, 15
FAN.CNT 1
                       KIND
                                        APPLICATION NO.
    PATENT NO.
                              DATE
                                                                DATE
                                         -----
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                                                                -----
     -----
                              -----
                              19941128 CA 1994-2123856
    CA 2123856
US 5455160
                       A1
                                                                19940518
PΙ
                                         US 1993-67802
                       Α
                              19951003
                                                               19930527
                       A
PRAI US 1993-67802
                              19930527
CLASS
 PATENT NO.
               CLASS PATENT FAMILY CLASSIFICATION CODES
 -----
               ____
 CA 2123856
               ICM
                       G01N033-68
                ICS
                       G01N033-574
                       G01N0033-68 [ICM,5]; G01N0033-574 [ICS,5]
                IPĊI
                       G01N0033-574 [I,C*]; G01N0033-574 [I,A]; G01N0033-68
                IPCR
                       [I,C*]; G01N0033-68 [I,A]
                       G01N0033-53 [ICM, 6]; G01N0033-574 [ICS, 6]
 US 5455160
                IPCI
                       G01N0033-574 [I,C*]; G01N0033-574 [I,A]; G01N0033-68
                IPCR
                       [I,C*]; G01N0033-68 [I,A]
                       435/007.230; 435/007.920; 435/007.930; 435/961.000;
                NCL
                       436/064.000; 436/811.000; 436/813.000
                ECLA
                       G01N033/574K; G01N033/68
AB
     This study describes a method for extraction and quantification of calprotectin
     (L1 protein) in feces by enzyme immunoassay. This protein is a prominent
     antimicrobial component of neutrophils, monocytes, macrophages and
     squamous epithelia. Calprotectin was stable in feces during storage for 7
     days at room temperature Fecal calprotectin quantitated in a 5-g sample taken
     from a 24-h feces collection gave a reliable estimate of calprotectin found in
     the pooled collection. The assay had a within assay precision (CV) of
     1.9% and a between assay precision of 14.8%. The following mean fecal
     calprotectin levels were found: healthy subjects 3095 µg/L; hospital
     controls 14,637 µg/L; and patients with inflammatory bowel disease
     (Crohn's disease and ulcerative colitis) 40,850 µg/L. The difference
     between the means are highly significant. All patients with IBD and 10 of
     11 patients with gastrointestinal carcinomas had calprotectin level above
     the suggested reference limit of 9000 µg/L. Determination of fecal
calprotectin is
     an important routine parameter for monitoring IBD and
     gastrointestinal cancer.
     feces calprotectin detn diagnosis disease cancer; enzyme immunoassay
ST
     calprotectin feces; inflammatory bowel disease diagnosis calprotectin;
     gastrointestinal tract calprotectin detn; colorectal carcinoma diagnosis
     calprotectin detn
IT
        (calprotectin determination in feces by EIA in diagnosis of disease or
       disorders of digestive system)
IT
     Antibodies
     RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); THU
     (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (calprotectin determination in feces by EIA in diagnosis of disease or
```

ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN

disorders of digestive system)

IT Bone marrow

(depression; calprotectin determination in feces by EIA in diagnosis of disease

or disorders of digestive system)

IT Intestine, disease

(Crohn's, calprotectin determination in feces by ${\tt EIA}$ in diagnosis of disease or

disorders of digestive system)

IT Proteins, specific or class

RL: ANT (Analyte); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); USES (Uses)

(L1, calprotectin determination in feces by EIA in diagnosis of disease or disorders of digestive system)

IT Digestive tract

(disease, calprotectin determination in feces by EIA in diagnosis of disease or

disorders of digestive system)

IT Intestine, disease

(inflammatory, calprotectin determination in feces by EIA in diagnosis of disease or disorders of digestive system)

IT Intestine, neoplasm

(large, carcinoma, calprotectin determination in feces by EIA in diagnosis

disease or disorders of digestive system)

IT Digestive tract

(neoplasm, carcinoma, calprotectin determination in feces by EIA in diagnosis

of disease or disorders of digestive system)

IT Intestine, disease

(ulcerative colitis, calprotectin determination in feces by EIA in diagnosis of

disease or disorders of digestive system)

οf

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ANSWER 1 OF 5 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN
     DUPLICATE 1
     2001:33185 BIOSIS
ΑN
DN
     PREV200100033185
    Natalizumab. Treatment of IBD, treatment of multiple
ΤI
     sclerosis: AN100226, AntegrenTM.
     Sorbera, L. A. [Reprint author]; Martin, L. [Reprint author]; Rabasseda,
ΑU
    X. [Reprint author]
CS
     Prous Science, 08080, Barcelona, Spain
    Drugs of the Future, (September, 2000) Vol. 25, No. 9, pp.
SO
     917-921. print.
     ISSN: 0377-8282.
DT
     Article
LA
    English
ED
     Entered STN: 10 Jan 2001
     Last Updated on STN: 12 Feb 2002
     Pharmacology - Clinical pharmacology
     Pathology - Therapy
                           12512
     Digestive system - Pathology
                                   14006
     Nervous system - Pathology 20506
     Pharmacology - General
                              22002
     Pharmacology - Immunological processes and allergy
                                                          22018
     Immunology - General and methods
                                       34502
     Immunology - Immunopathology, tissue immunology ·34508
IT
     Major Concepts
        Clinical Immunology (Human Medicine, Medical Sciences);
        Gastroenterology (Human Medicine, Medical Sciences); Pharmacology
IT
    Diseases
        inflammatory bowel disease: digestive system disease
        Inflammatory Bowel Diseases (MeSH)
IT
     Diseases
        multiple sclerosis: immune system disease, nervous system disease
        Multiple Sclerosis (MeSH)
IT
     Chemicals & Biochemicals
        natalizumab [AN 100226, Antegren]: immunosuppressant-drug, monoclonal
        antibody
ORGN Classifier
        Hominidae
                    86215
     Super Taxa
        Primates; Mammalia; Vertebrata; Chordata; Animalia
     Organism Name
        human
     Taxa Notes
        Animals, Chordates, Humans, Mammals, Primates, Vertebrates
     189261-10-7 (natalizumab)
     189261-10-7 (AN 100226)
     189261-10-7 (Antegren)
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10/629,975 Search LVCOOK 6/25/07

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(FILE 'HOME' ENTERED AT 17:25:47 ON 27 JUN 2007)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 17:26:06 ON 27 JUN 2007

	JUN 2007						
Ll	66455	S	(ULCERAT	TIVE COL	ITIS)		
L2	248	S	L1 AND I	LACTOFER	RIN?		
L3	23	S	L2 AND T	REATMEN'	r?		
L4	17	DI	JPLICATE	REMOVE :	L3 (6	DUPLICATES	REMOVED)
L5	7	S	L4 AND E	PD<2001	•		
L6	35	S	L2 AND T	THERAPY			
L7	35	S	L6 NOT I	1 5			
L8	27	DI	JPLICATE	REMOVE :	L7 (8	DUPLICATES	REMOVED)
L9	9	S	L8 AND B	PD<2001			

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(FILE 'HOME' ENTERED AT 17:25:47 ON 27 JUN 2007)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 17:26:06 ON 27 JUN 2007

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L2	248	S	L1 AN	D LAC	COFE	RRIN	1?		
L3	23	S	L2 AN	D TREA	ATME	YT?			
L4	17	ĎĮ	JPLICA	TE RE	OVE	L3	(6	DUPLICATES	REMOVED)
L5	· 7	S	L4 AN	D PD<2	2001				
L6	35	S	L2 AN	D THE	RAPY				
L7	. 35	S	L6 NO	T L5					
L8	27	D	JPLICA	TE RE	40VE	L7	(8	DUPLICATES	REMOVED)
T.Q	۵	C	T.O 7\N	י-חם ח	2001				•

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ANSWER 6 OF 9 EMBASE COPYRIGHT (c) 2007 Elsevier B.V. All rights
     reserved on STN
     1999322801 EMBASE
ΑN
TI
     Faecal parameters in the assessment of activity in inflammatory bowel
AU
     Van der Sluys Veer A.; Biemond I.; Verspaget H.W.; Lamers C.B.H.W.
     A. Van der Sluys Veer, Dept of Gastroenterol. and Hepatol., Leiden
     University Medical Center, Building 1, PO Box 9600, C4-PNL-2300 RC Leiden,
     Netherlands
SO
     Scandinavian Journal of Gastroenterology, Supplement, (1999)
     Vol. 33, No. 230, pp. 106-110. .
     Refs: 55
     ISSN: 0085-5928 CODEN: SJGSB8
CY
     Norway
     Journal; Article
DT
FS
     005
             General Pathology and Pathological Anatomy
     029
             Clinical Biochemistry
     037
             Drug Literature Index
     048
             Gastroenterology
     English
LA
SL
     English
ED
     Entered STN: 30 Sep 1999
     Last Updated on STN: 30 Sep 1999
AB
     Background: Determination of inflammatory activity is helpful when
     assessing the efficacy of drugs in therapeutic trials and in facilitating
     management of individual patients with inflammatory bowel disease (IBD).
     Faecal parameters have been hypothesized to be more specific than
     non-faecal measurements in the assessment of intestinal inflammation.
     Methods: Review of the literature on faecal measurements in IBD. Results
     and conclusions: Leakage of various proteins and leukocyte products into
     the intestinal lumen can be assessed and quantified in stool specimens and
     serve as a measurement of inflammatory activity. Several of these faecal
     parameters are raised in patients with IBD. There is a considerable
     overlap between patients with active and those with inactive disease,
     however, and the correlation of the faecal parameters with disease
     activity indices is often low. The value of \alpha.apprx.1-antitrypsin
     measurement in faeces in the assessment of intestinal inflammation has
     been well established. Further studies in patients with IBD are needed to
     determine whether other faecal parameters, such as lactoferrin,
     tumour necrosis factor \alpha, PMN-elastase, lysozyme, leucocyte
     esterase, immunoglobulin A, among others, are more accurate or
     cost-effective than measurement of \alpha.apprx.1-antitrypsin in the
     stools of such patients.
    Medical Descriptors:
     *feces
     *enteritis: DI, diagnosis
       *enteritis: DT, drug therapy
     *Crohn disease: DI, diagnosis
       *Crohn disease: DT, drug therapy
       *ulcerative colitis: DI, diagnosis
       *ulcerative colitis: DT, drug therapy
     gastrointestinal endoscopy
     intestine biopsy
     imaging
     immunodiffusion
     enzyme immunoassay
     nephelometry
     human
     clinical trial
     article
     priority journal
     Drug Descriptors:
     *alpha 1 antitrypsin: EC, endogenous compound
     protein: EC, endogenous compound
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ANSWER 6 OF 9 EMBASE COPYRIGHT (c) 2007 Elsevier B.V. All rights
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     1999322801 EMBASE
AN
ΤI
     Faecal parameters in the assessment of activity in inflammatory bowel
ΑU
     Van der Sluys Veer A.; Biemond I.; Verspaget H.W.; Lamers C.B.H.W.
     A. Van der Sluys Veer, Dept of Gastroenterol. and Hepatol., Leiden
     University Medical Center, Building 1, PO Box 9600, C4-PNL-2300 RC Leiden,
     Netherlands
SO
     Scandinavian Journal of Gastroenterology, Supplement, (1999)
     Vol. 33, No. 230, pp. 106-110. .
     Refs: 55
     ISSN: 0085-5928 CODEN: SJGSB8
CY
     Norway
DT
     Journal; Article
FS
     005
             General Pathology and Pathological Anatomy
     029
             Clinical Biochemistry
     037
             Drug Literature Index
     048
             Gastroenterology
LΑ
     English
SL
     English
ED
     Entered STN: 30 Sep 1999
     Last Updated on STN: 30 Sep 1999
AB
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     esterase, immunoglobulin A, among others, are more accurate or
     cost-effective than measurement of \alpha.apprx.1-antitrypsin in the
     stools of such patients.
CT
     Medical Descriptors:
     *feces
     *enteritis: DI, diagnosis
       *enteritis: DT, drug therapy
     *Crohn disease: DI, diagnosis
       *Crohn disease: DT, drug therapy
       *ulcerative colitis: DI, diagnosis
       *ulcerative colitis: DT, drug therapy
     gastrointestinal endoscopy
     intestine biopsy
     imaging
     immunodiffusion
     enzyme immunoassay
     nephelometry
     human
     clinical trial
     article
     priority journal
     Drug Descriptors:
     *alpha 1 antitrypsin: EC, endogenous compound
     protein: EC, endogenous compound
```

```
lactoferrin: EC, endogenous compound
     tumor necrosis factor alpha: EC, endogenous compound
     leukocyte elastase: EC, endogenous compound
     lysozyme: EC, endogenous compound esterase: EC, endogenous compound
     immunoglobulin a: EC, endogenous compound
     barium
     methylprednisolone: CB, drug combination
       methylprednisolone: DT, drug therapy
     salazosulfapyridine: CB, drug combination
       salazosulfapyridine: DT, drug therapy
     hemoglobin: EC, endogenous compound
     indium 111
     (alpha 1 antitrypsin) 9041-92-3; (protein) 67254-75-5; (
RN
     lactoferrin) 55599-62-7; (leukocyte elastase) 109968-22-1;
     (lysozyme) 9001-63-2; (esterase) 9013-79-0; (barium) 7440-39-3;
     (methylprednisolone) 6923-42-8, 83-43-2; (salazosulfapyridine) 599-79-1;
     (hemoglobin) 9008-02-0; (indium 111) 15750-15-9
```

```
lactoferrin: EC, endogenous compound
     tumor necrosis factor alpha: EC, endogenous compound
     leukocyte elastase: EC, endogenous compound
     lysozyme: EC, endogenous compound
     esterase: EC, endogenous compound
     immunoglobulin a: EC, endogenous compound
     barium
     methylprednisolone: CB, drug combination
       methylprednisolone: DT, drug therapy
     salazosulfapyridine: CB, drug combination
       salazosulfapyridine: DT, drug therapy
     hemoglobin: EC, endogenous compound
     indium 111
RN
     (alpha 1 antitrypsin) 9041-92-3; (protein) 67254-75-5; (
     lactoferrin) 55599-62-7; (leukocyte elastase) 109968-22-1;
     (lysozyme) 9001-63-2; (esterase) 9013-79-0; (barium) 7440-39-3;
     (methylprednisolone) 6923-42-8, 83-43-2; (salazosulfapyridine) 599-79-1;
     (hemoglobin) 9008-02-0; (indium 111) 15750-15-9
```

ANSWER 4 OF 9 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN AN 1993:396686 BIOSIS DN PREV199345055511 TI A study to determine fecal lactoferrin in patients with ulcerative colitis. ΑU Sudo, Ichiro CS Fourth Dep. Intern. Med., Tokyo Med. Coll., Japan SO Japanese Journal of Gastroenterology, (1993) Vol. 90, No. 4, pp. 824. ISSN: 0446-6586. DT Article LΑ Japanese Entered STN: 30 Aug 1993 ED Last Updated on STN: 30 Aug 1993 CC Biochemistry studies - Proteins, peptides and amino acids 10064 Pathology - Diagnostic 12504 Pathology - Inflammation and inflammatory disease Pathology - Therapy 12512 Digestive system - General and methods Digestive system - Pathology Immunology - Immunopathology, tissue immunology IT Major Concepts Clinical Endocrinology (Human Medicine, Medical Sciences); Digestive System (Ingestion and Assimilation); Gastroenterology (Human Medicine, Medical Sciences); Pathology Miscellaneous Descriptors IT DIAGNOSTIC METHOD; PATHOLOGY; THERAPY ORGN Classifier 86215 Hominidae Super Taxa Primates; Mammalia; Vertebrata; Chordata; Animalia Organism Name human Taxa Notes

Animals, Chordates, Humans, Mammals, Primates, Vertebrates

ANSWER 4 OF 9 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN AN 1993:396686 BIOSIS DN PREV199345055511 A study to determine fecal lactoferrin in patients with TI ulcerative colitis. Sudo, Ichiro ΑU Fourth Dep. Intern. Med., Tokyo Med. Coll., Japan CS Japanese Journal of Gastroenterology, (1993) Vol. 90, No. 4, pp. SO 824. ISSN: 0446-6586. DTArticle LΑ Japanese Entered STN: 30 Aug 1993 ED Last Updated on STN: 30 Aug 1993 CC Biochemistry studies - Proteins, peptides and amino acids 10064 Pathology - Diagnostic 12504 Pathology - Inflammation and inflammatory disease Pathology - Therapy 12512 Digestive system - General and methods 14001 Digestive system - Pathology Immunology - Immunopathology, tissue immunology IT Major Concepts Clinical Endocrinology (Human Medicine, Medical Sciences); Digestive System (Ingestion and Assimilation); Gastroenterology (Human Medicine, Medical Sciences); Pathology IT Miscellaneous Descriptors DIAGNOSTIC METHOD; PATHOLOGY; THERAPY ORGN Classifier Hominidae 86215 Super Taxa Primates; Mammalia; Vertebrata; Chordata; Animalia human Taxa Notes

Animals, Chordates, Humans, Mammals, Primates, Vertebrates

```
ANSWER 1 OF 9 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN
     1994:286260 BIOSIS
AN
     PREV199497299260
DN
     Mucosal lactoferrin, "sticky" neutrophils, and pathergy in
TI
     ulcerative colitis and pyoderma gangrenosum:
     Implications for pathogenesis, and successful therapy with
ΑU
     Dwarakanath, A. D. [Reprint author]; Finnie, I. A. [Reprint author]; Yu,
     L. G. [Reprint author]; O'Dowd, G. M.; Rhodes, Jonathan M.
CS
     Dep. Med., Univ. Liverpool, P.O. Box 147, Liverpool L69 3BX, UK
     Gastroenterology, (1994) Vol. 106, No. 4 SUPPL., pp. A674.
SO
     Meeting Info.: 95th Annual Meeting of the American Gastroenterological
     Association. New Orleans, Louisiana, USA. May 15-18, 1994.
     CODEN: GASTAB. ISSN: 0016-5085.
DT
     Conference; (Meeting)
     Conference; Abstract; (Meeting Abstract)
LA
     English
ED
     Entered STN: 30 Jun 1994
     Last Updated on STN: 1 Jul 1994
CC
     General biology - Symposia, transactions and proceedings
     Cytology - Human
                        02508
     Biochemistry studies - Carbohydrates
                                            10068
     Biophysics - Molecular properties and macromolecules
     Pathology - Inflammation and inflammatory disease
     Pathology - Therapy
                           12512
     Digestive system - Pathology
                                    14006
     Blood - Blood cell studies
                                  15004
     Blood - Blood, lymphatic and reticuloendothelial pathologies
     Blood - Lymphatic tissue and reticuloendothelial system
     Pharmacology - Clinical pharmacology
     Pharmacology - Digestive system
IT
     Major Concepts
        Biochemistry and Molecular Biophysics; Blood and Lymphatics (Transport
        and Circulation); Cell Biology; Gastroenterology (Human Medicine,
        Medical Sciences); Hematology (Human Medicine, Medical Sciences);
        Pathology; Pharmacology
     Chemicals & Biochemicals
IT
        HEPARIN
IT
     Miscellaneous Descriptors
        GASTROINTESTINAL-DRUG; HEPARIN; MEETING ABSTRACT
ORGN Classifier
        Hominidae
                    86215
     Super Taxa
        Primates; Mammalia; Vertebrata; Chordata; Animalia
     Organism Name
       human
     Taxa Notes
        Animals, Chordates, Humans, Mammals, Primates, Vertebrates
RN
     9005-49-6 (HEPARIN)
```

New/Noteworthy

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	Limits	, Preview/Index	•	Clipboard	Details				
About Entrez	Display	itation		Show 20	Sort by	▼ Se	end to	▼	
Text Version	All: 1 F	Review: 0							•
Entrez PubMed Overview Help FAQ	EI	n Biochem. 199 LSEVIER L-TEXT ARTICLE	94 Aug;27(4):259-64.				Related A	rticles, Links

Uchida K, Matsuse R, Tomita S, Sugi K, Saitoh O, Ohshiba S.

inflammatory gastrointestinal disorders and colon cancer.

Kyoto Medical Science Laboratory, Kyoto, Japan.

We have developed a new immunochemical test for fecal lactoferrin (LF) utilizing an enzyme-linked immunosorbent assay (ELISA). The ELISA had a sensitivity of about 10 micrograms/L of lactoferrin and the measurable range was 10.0-1000.0 micrograms/L (1.0-100.0 micrograms LF/g feces). The stability of lactoferrin in feces was greater than that of myeloperoxidase and leucocyte elastase. The fecal concentration of lactoferrin (mean +/- SD) in 35 normal subjects was 0.75 +/- 0.83 microgram/g feces, whereas that in 24 patients with colon cancer was 74.4 +/- 88.3 micrograms/g feces. The fecal lactoferrin concentration of 38 patient with active ulcerative colitis was 307.4 +/- 233.9 micrograms/g feces, and that in 36 patients with active Crohn's disease was 191.7 +/- 231.1 micrograms/g feces. The ELISA for human fecal lactoferrin might be useful in the diagnosis of colon disease.

Immunochemical detection of human lactoferrin in feces as a new marker for

Publication Types:

• Comparative Study

MeSH Terms:

- Adolescent
- Adult
- Aged
- Colitis, Ulcerative/diagnosis
- Colitis, Ulcerative/metabolism*
- Colonic Neoplasms/diagnosis
- Colonic Neoplasms/metabolism*
- Colonic Polyps/diagnosis
- Colonic Polyps/metabolism
- Colonoscopy
- Crohn Disease/diagnosis
- Crohn Disease/metabolism*
- Enzyme-Linked Immunosorbent Assay
- Feces/chemistry*
- Female
- Humans
- Lactoferrin/blood

- Lactoferrin/metabolism*
- Male
- Middle Aged
- Regression Analysis

Substances:

• Lactoferrin

PMID: 8001286 [PubMed - indexed for MEDLINE]

Display Citation	•	▼ SI	how 20	▼ Sort by	▼ Send to ▼

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